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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/015,930	11/30/2001	Jane Hirsh	CP 104	2912

7590

01/31/2003

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EXAMINER

TRAN, SUSAN T

ART UNIT

PAPER NUMBER

1615

DATE MAILED: 01/31/2003

Please find below and/or attached an Office communication concerning this application or proceeding.

<b>Office Action Summary</b>	<b>Application No.</b> 10/015,930	<b>Applicant(s)</b> HIRSH ET AL.	
	<b>Examiner</b> Susan Tran	<b>Art Unit</b> 1615	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☐ Responsive to communication(s) filed on \_\_\_\_.
- 2a) ☐ This action is **FINAL**.                      2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 1-23 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 1-23 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
     Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on \_\_\_\_ is: a) ☐ approved b) ☐ disapproved by the Examiner.  
     If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

**Priority under 35 U.S.C. §§ 119 and 120**

- 13) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).  
     a) ☐ All    b) ☐ Some \* c) ☐ None of:  
         1. ☐ Certified copies of the priority documents have been received.  
         2. ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_.  
         3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).  
     \* See the attached detailed Office action for a list of the certified copies not received.
- 14) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).  
     a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

**Attachment(s)**

- |  |   |
|--|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892)                                  | 4) <input type="checkbox"/> Interview Summary (PTO-413) Paper No(s). ____.  |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948)                         | 5) <input type="checkbox"/> Notice of Informal Patent Application (PTO-152) |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO-1449) Paper No(s) <u>Z</u> . | 6) <input type="checkbox"/> Other: _____                                    |

### **DETAILED ACTION**

Receipt is acknowledged of applicant's Corrected Filing Receipt filed 01/24/02, Fee and Declaration filed 01/31/02, Power of Attorney filed 09/20/02, and Information Disclosure Statement filed 10/23/02.

### ***Claim Rejections - 35 USC § 103***

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

Claims 1-20, 22, and 23 are rejected under 35 U.S.C. 103(a) as being unpatentable over Conley et al. US 6,294,199.

Conley teaches method of treating a bacterial infection comprising administering composition comprises amoxycillin (see abstract). The composition can be a modified release dosage formulation comprises an immediate release phase and a slow release phase (column 9, lines 5-58). The modified release dosage form can be a dispersible tablet, a swallow tablet, a chewable tablet that may also be effervescent, a capsule or a sachet (column 9, lines 66 through column 10, lines 1-3). The modified release tablet can be formulated in a bi-layered tablet comprises an immediate release layer which comprises amoxycillin, disintegrants, compression aids, diluents, lubricants, and the like, which will disintegrate rapidly; and a slow release layer which comprises

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amoxicillin together with release retarding polymers (columns 11-12). Columns 15-17 disclose the process of making the composition.

Conley is silent as to the teaching that the immediate release layer dissolved intraorally. However, Conley teaches the modified release formulation that can be formulated in a chewable tablet. Thus, such language does suggest the active agent in the immediate release layer disintegrates rapidly in the mouth, and therefore, provide intraoral absorption. Accordingly, it would have been *prima facie* obvious for one of ordinary skill in this art to optimize Conley's modified release formulation with the expectation of at least similar result, because Conley teaches the advantageous result in the use of bi-layer chewable/effervescent tablet comprising active agent in both, the immediate release layer, and the delay/slow release layer.

Claim 21 is rejected under 35 U.S.C. 103(a) as being unpatentable over Conley et al.

Conley is relied upon for the reason stated above. The reference is silent as to the method of administering the modified release formulation as claimed in claim 21. However, Conley teaches a bi-layer chewable tablet that comprises an immediate release layer that will disintegrate rapidly; and a slow release layer which after oral ingestion will disintegrate/dissolve in the intestine (columns 11-12). Thus, it is the position of the examiner that such language does suggests that after the effervescent of the immediate release layer in the mouth, the ingestion of the slow release layer then occurs. Therefore, it would have been *prima facie* obvious for one of ordinary skill in the

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art to, by routine experimentation determine a suitable administration method with the expectation of at least similar result, because Conley teaches a modify release formulation having biphasic release profile that fall within the claimed range (column 9, lines 28-44).

Claim 1 is rejected under 35 U.S.C. 103(a) as being unpatentable over Van Scoik US 5,082,667, and Hackhs (Chemical Dictionary).

Van Scoik teaches a pharmaceutical formulation in tablet triturate form comprising carbohydrate and active agent particle that release active ingredient after digestion (columns 2-4).

Van Scoik does not teach active agent which disintegrates/dissolves during the quick dissolution of the triturate tablet. However, it is the position of the examiner that the generic term "carbohydrate" of Van Scoik would suggest the use of active agent.

Hackhs teaches carbohydrates include polysaccharides, which includes starches, cellulose, glycogens, and inulins (pages 185-186). Thus, it would have been obvious for one of ordinary skill in the art to modify Van Scoik to incorporate an active agent in the quick dissolve portion of the triturate tablet in view of the teaching of Hackhs to obtain the claimed invention, because Van Scoik teaches the advantageous results in the use of triturate tablet containing carbohydrates useful in pharmaceutical art.

***Pertinent Arts***

The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. Wilen, Fusari et al., Schmitt, Porter, Gergely et al., Van Scoik, and Dsai are cited as being of interest for the teachings of triturate and/or effervescent tablet.

***Correspondence***

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Susan Tran whose telephone number is (703) 306-5816. The examiner can normally be reached on Monday through Thursday from 6:00 am to 4:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Thurman Page, can be reached on (703) 308-2927. The fax phone number for the organization where this application or proceeding is assigned is (703) 305-3592.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-1235.

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